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(se) Quinoline derivatives, processes for their preparation, their use, pharmaceutical compositions containing them and a method for the preparation of these pharmaceutical compositions.

The invention is concerned with novel quinicine and cinchonicine derivatives having cardiovascular activities of the formula or a salt thereof,

$$\begin{array}{c} A-B-CH_2 \\ R^1 \\ \end{array}$$

in which A-B is -CH₂-CH₂, -CHOH-CH₂, -CH₂-CHOH-,-C(O)-CH₂, -CH₂-C(O)-, -C(NOR*)-CH₂-or -CH₂-C(NOR*)-; R¹ is hydrogen, hydroxy or lower alkoxy; R³ is ethyl or vinyl; R² is Caa alkyl, C1-a hydroxyalkyl, lower alkoxy-alkyl or tower alkoxy-cycloalkyl, cower alkoxy-cycloalkyl, bower alkoxy-cycloalkyl or lower alkoxy-or lower alkyl, hydroxy-, lower alkoxy- or lower alkyl, cycloalkyl lower alkyl; cyano, cyano lower alkyl, lower alkoyl, lower alkyl, tetrahydrofuryl, mono- or di-lower alkylamino

lower alkyl, mono- or di-lower alkylamino lower hydroxy alkyl; optionally substituted phenyl, phenyl lower alkyl or phenyl hydroxy lower alkyl, optionally substituted diphenyl lower alkyl, optionally substituted phenyl lower alkyl, optionally substituted benzoyl or benzoyl lower alkyl, or optionally substituted heteroaryl or heteroaryl lower alkyl, or optionally substituted heteroaryl or heteroaryl lower alkyl, or optionally substituted heteroaryl or heteroaryl lower alkyl, R⁴ is lower alkyl, and Z is hydrogen, lower alkyl or optionally substituted phenyl, or Z and R³ together with the carbon atom to which they are attached form a Cas cycloalkyl group, whereby the substitutents at the 3- and 4-position of the piperidine ring are in the cis-position, excluding N-[Cas alkyl, optionally substituted Cr-11 aralkyl] substituted derivatives of quinicine and cinchonicine.

The compounds of the formula may be in the form of their optically active enantiomers and/or their therapeutically acceptable salts.

Methods for the preparation of the compounds of the formula are also disclosed and form part of the invention.